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UNITED STATES ENVIRONMENTAL PROTECTION AGENCY WASHINGTON, D.C. 20460

MAR 1 9 1987

MEMORANDUM

OFFICE OF
PESTICIDES AND TOXIC SUBSTANCE

Addition of new impurity to bifenthrin technical Subject:

To: George La Rocca (PM-15)

Registration Division, TS-767C

From: Marcia van Gemert, Ph.D. W. waw June 3/18/87
Head, Section III
Toxicology Branch, HED

Chief, Toxicology Branch, HED

Chemical: Bifenthrin, Talstar, FMC 54800

Project No: 7-0484

Caswell No: 763F

EPA ID NO.: 279-3055

Action Requested: Registration Division has sent a request to Toxicology Branch to determine if additional toxicology data are necessary to approve this formulation change.

Comments: FMC has submitted information concerning their new manufacturing process that introduces a new impurity into the final technical bifenthrin. TFP acid, the starting material for bifenthrin, is the compound

(referred to as IS acid impurity). During the manuracturing process TFP acid is esterified. addition this new impurity is esterified into

(referred to as TS bifenthrin

impurity).

This TS acid impurity can be present at levels as high as FMC claims that this impurity cannot be realistically reduced below about below about FMC also states that a high percent of the impurity present is probably transferred into the ester impurity in the Bifenthrin technical.

CARLES AND PROCESS INTORIALIZATION IS NOT COLUMN.

Based on the fact that a significant amount of this new impurity will be coming into the technical bifenthrin, and all the toxicology studies done to date have not incorporated this new impurity into their testing, the Toxicology Branch will require that an acute oral toxicity study and a full mutagenicity battery (similar to that submitted for the original bifenthrin technical for comparison) be performed on this new TS bifenthrin impurity and submitted for our review before this new formulation can replace the old one.

Structures of bifenthrin, TS acid impurity and TS bifenthrin impurity are on appended page 1.

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	Identity of product impurities.
	Description of the product manufacturing process.
	Description of quality control procedures.
	Identity of the source of product ingredients.
	Sales or other commercial/financial information.
	A draft product label.
	The product confidential statement of formula.
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